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<p>(21) International Application Number: PCT/EP96/02961</p> <p>(22) International Filing Date: 5 July 1996 (05.07.96)</p> <p>(30) Priority Data: 9515412.6 27 July 1995 (27.07.95) GB</p> <p>(71) Applicant (<i>for all designated States except US</i>): PHARMACIA & UPJOHN S.P.A. [IT/IT]; Via Robert Koch, 1,2, I-20152 Milan (IT).</p> <p>(72) Inventors and (75) Inventors/Applicants (<i>for US only</i>): PEVARELLO, Paolo [IT/IT]; Piazza San Pietro in Ciel d'Oro, 7/A, I-27100 Pavia (IT). AMICI, Raffaella [IT/IT]; Via N. Rocca, 11, I-29100 Piacenza (IT). VARASI, Mario [IT/IT]; Via Giambellino, 80, I-20146 Milan (IT). BONSIGNORE, Alberto [IT/IT]; Via dei Benedettini, 2, I-20146 Milan (IT). SALVATI, Patricia [IT/IT]; Via Valeca, 16/C, I-20020 Arese (IT).</p>		<p>(81) Designated States: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CZ, DE, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, RO, RU, SD, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).</p> <p>Published <i>With international search report.</i></p>	
<p>(54) Title: 2-(4-SUBSTITUTED)-BENZYLAMINO-2-METHYL-PROPANAMIDE DERIVATIVES</p> <p style="text-align: center;"> (I) </p> <p>(57) Abstract</p> <p>Novel 2-(4-substituted)-benzylamino-2-methyl-propanamides compounds, having CNS activity, of general formula (I) wherein: n is zero, 1, 2 or 3; X is -O-, -S-, -CH₂- or -NH-; each of R and R₁ independently is hydrogen, C₁-C₆ alkyl, halogen, hydroxy, C₁-C₆ alkoxy or trifluoromethyl; each of R₂, R₃ and R₄ independently is hydrogen, C₁-C₆ alkyl or C₃-C₆ cycloalkyl; and the pharmaceutically acceptable salts thereof, are disclosed.</p>			

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**2-(4-SUBSTITUTED)-BENZYLAMINO-2-METHYL-PROPANAMIDE
DERIVATIVES**

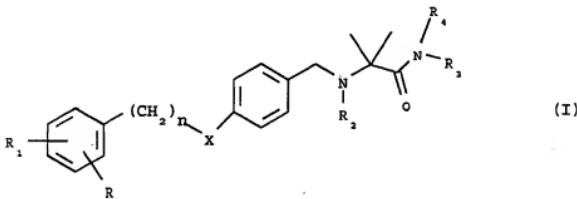
The present invention relates to novel 2-(4-substituted)-benzylamino-2-methyl-propanamides, to their use as therapeutic agents, to a process for their preparation and to pharmaceutically compositions containing them.

WO 90/14334 discloses N-phenylalkyl substituted α -amino carboxamide derivatives active on the CNS.

It has been found that novel 2-(4-substituted)-benzylamino-2-methyl-propanamides derivatives as herein defined have valuable biological properties, in particular as antiepileptic, anti-Parkinson, neuroprotective, anti-depressant, antispastic and/or hypnotic agent.

15

The present invention provides novel compounds of the following general formula (I)



wherein:

- 20 n is zero, 1, 2 or 3;
- X is -O-, -S-, -CH₂- or -NH-;
- each of R and R₁ independently is hydrogen, C₁-C₆ alkyl, halogen, hydroxy, C₁-C₄ alkoxy or trifluoromethyl;
- each of R₂, R₃ and R₄ independently is hydrogen, C₁-C₆ alkyl or C₃-C₇ cycloalkyl;
- and the pharmaceutically acceptable salts thereof.

The pharmaceutically acceptable salts of the compounds of formula (I) include acid addition salts with inorganic, e.g. hydrochloric, hydrobromic, sulfuric, and phosphoric acids, or organic, e.g. acetic, propionic, lactic, oxalic, malic, 5 maleic, tartaric, citric, benzoic, mandelic, C₁-C₄ alkylsulfonic, salicylic and fumaric acids.

The compounds of the formula (I), their pharmaceutically acceptable salts may also form pharmaceutically acceptable solvates, such as mono-, di- or tri-hydrates, which are also 10 object of the present invention.

The alkyl and alkoxy groups may be branched or straight groups.

A C₁-C₆ alkyl group is preferably a C₁-C₄ alkyl group, in particular methyl, ethyl, n- and iso-propyl, n-, iso-, sec- 15 and tert-butyl, more preferably methyl or ethyl.

Representative examples of C₁-C₄ alkoxy groups include methoxy or ethoxy.

A halogen atom is e.g. chlorine, fluorine or bromine.

A C₃-C₇ cycloalkyl group is, for instance, a cyclopropyl, 20 cyclohexyl or cycloheptyl group, in particular cyclopropyl.

The present invention also include within its scope all the possible isomers of the compounds of formula (I) and their mixtures, as well as the metabolites and the pharmaceutically acceptable bio-preursors (otherwise known as pro-drugs) of 25 the compounds of formula (I).

Preferred compounds of the invention are the compounds of formula (I) wherein

n is 1 or 2;

30 X is -O-, -S- or -NH-;

R is hydrogen;

R₁ is hydrogen or halogen;

each of R₂, R₃ and R₄ independently is hydrogen or C₁-C₄ alkyl; and the pharmaceutically acceptable salts thereof.

More preferred compounds of the invention are the compounds
5 of formula (I), wherein

n is 1;

X is -O-, -S- or -NH-;

R₁ is hydrogen or halogen;

R, R₂, R₃ and R₄ are hydrogen; and the pharmaceutically
10 acceptable salts thereof.

Examples of specific compounds of the invention are:

2-[4-(3-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;

2-[4-(3-chlorobenzyl)oxy]benzylamino]-2-methyl-propanamide;

15 2-[4-(4-chlorobenzyl)oxy]benzylamino]-2-methyl-propanamide;

2-[4-(3-bromobenzyl)oxy]benzylamino]-2-methyl-propanamide;

2-[4-(4-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;

2-[4-(2-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;

2-[4-(3-fluorobenzyl)amino]benzylamino]-2-methyl-propanamide;

20 2-[4-(benzylsulfanyl)benzylamino]-2-methyl-propanamide;

2-[4-(3-fluorobenzyl)oxy]benzylmethylamino]-2-methyl-
propanamide;

2-[4-(3-fluorobenzyl)oxy]benzyl-amino]-2-methyl-N-methyl-
propanamide;

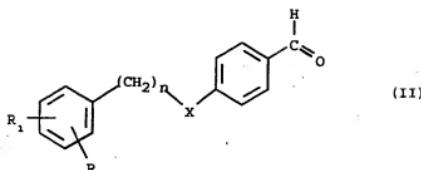
25 2-[4-(3-chlorobenzyl)oxy]benzylmethylamino]-2-methyl-
propanamide; and

2-[4-(3-bromobenzyl)oxy]benzylmethylamino]-2-methyl-
propanamide, if the case, either as single (S) or (R) isomer
or as a mixture thereof, and the pharmaceutically acceptable
30 salts thereof.

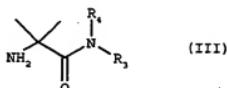
The compounds of the invention and the salts thereof can be

obtained by a process comprising:

- a) reacting a compound of formula (II)



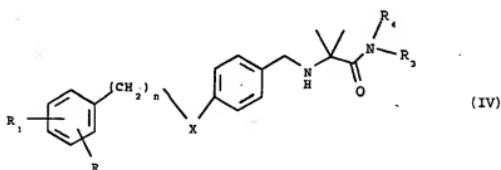
5 wherein n, R, R₁ and X are as defined above, with a compound of formula (III)



wherein R₃ and R₄ are as defined above, thus obtaining a compound of formula (I) in which R₂ is hydrogen; or

10

- b) reacting a compound of formula (IV)



wherein R, R₁, R₃, R₄, n and X are as defined above, with a compound of formula (V) or (VI)

15



wherein W is a halogen atom; R'₂ is a C₁-C₆ alkyl and R''₂ is hydrogen or C₁-C₅ alkyl, thus obtaining a compound of the invention in which R₂ is C₁-C₆ alkyl; and, if desired, converting a compound of the invention into another compound of the invention and/or, if desired, converting a compound of the invention into a

20

pharmaceutically acceptable salt and/or, if desired, converting a salt into a free compound.

All the processes described hereabove are analogy processes
5 and can be carried out according to well known methods in organic chemistry.

A compound of formula (IV) is a compound of formula (I) in which R₂ is hydrogen.

The reaction of a compound of formula (II) with a compound of formula (III) to give a compound of formula (I) or (IV) is a reductive amination reaction which can be carried out according to well known methods. According to a preferred embodiment of the invention it may be performed under nitrogen atmosphere, in a suitable organic solvent, such as
10 an alcohol, e.g. a lower alkanol, in particular methanol, or in acetonitrile, at a temperature ranging from about 0°C to about 40°C, in the presence of a reducing agent, the most appropriate being sodium cyanoborohydride. Occasionally molecular sieves can be added to the reaction mixture for
15 facilitating the reaction.

In a compound of formula (V) the halogen W is preferably iodine. The alkylation reaction of a compound of formula (IV) with a compound of formula (V) can be carried out in a suitable organic solvent, such as an alcohol, e.g. methanol,
20 ethanol or isopropanol, in particular in ethanol, at a temperature ranging from about 0°C to about 50°C.

The alkylation reaction of a compound of formula (IV) with an aldehyde of formula (VI) can be carried out in a suitable organic solvent, such as an alcohol, e.g. methanol, ethanol or acetonitrile in the presence of a suitable reducing agent, such as sodium cyanoborohydride, at a temperature ranging
30 from about 0°C to about 30°C.

A compound of the invention can be converted, as stated above, into another compound of the invention by known methods. Process-variant b) above may be regarded as an example of optional conversion of a compound of the invention
5 into another compound of the invention.

Also the optional salification of a compound of the invention as well as the conversion of a salt into the free compound may be carried out by conventional methods.

The compounds of formula (II), (III), (V) and (VI) are known
10 compounds or can be obtained by known methods.

When in the compounds of the present invention and in the intermediate-products thereof, groups are present, which need to be protected before submitting them to the hereabove illustrated reactions, they may be protected before being
15 reacted and then deprotected according to methods well known in organic chemistry.

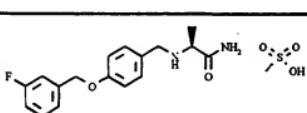
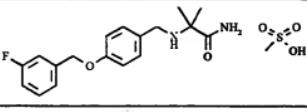
PHARMACOLOGY

The compounds of the invention are active on the central nervous system (CNS) and can be used in therapy, for example as antiepileptics, in the treatment of Parkinson's disease and as neuroprotective agents, e.g. preventing or treating neuronal loss associated with stroke, hypoxia, ischemia, CNS trauma, hypoglycaemia or surgery and in treating and
25 preventing neurodegenerative diseases such as Alzheimer's disease, amyotrophic lateral sclerosis, Down's syndrome, Huntington's disease, dementia caused by acquired immunodeficiency syndrome (AIDS), infarctual dementia and infections or inflammations in the brain; they can also be used as antidepressants, hypnotics and antispastic agents and
30 in treating ocular damage and retinopathy.

The activity on the CNS of the compounds of the invention was

evaluated on the basis of pharmacological methods, such as, for example, the antagonism of convulsions and lethality induced by intravenous injection of bicuculline in mice (Antiepileptic Drugs, D.M. Woodbury et al. eds., 2nd edition, 5 Raven Press, New York, 1982), or the antagonism of maximal electroshock seizures (MES) (Woodbury, L.A. and Davenport, V.D., Arch. Int. Pharmacodyn. Ther. 92, 97-104, 1952).
 The following table summarizes the activity data obtained by testing in the MES test a representative group of compounds 10 of the invention in comparison with the respective 2-desmethyl related compounds known from WO 90/14334.
 The symbol * identifies the prior art 2-desmethyl-compounds known from WO 90/14334.
 ED₅₀ means effective dose in 50% of treated animals after per 15 os (po) administration.

Table 1

Structure	Compound (internal code)	MES-ED ₅₀ (po) mg/Kg
	FCE 26743A*	8.2 (7.2-9.4)
	FCE 29088A	4.4 (2.8-6.3)

(cont..)

Table 1 (cont'd)

Structure	Compound (internal code)	MES-ED ₅₀ (po) mg/Kg
	FCE 26727A*	12.72 (9.19-17.62)
	FCE 29482A	3.12 (2.28-4.27)
	FCE 26742A*	7.13 (5.70-8.91)
	FCE 29484A	3.10 (2.25-4.28)
	FCE 26193A*	9.41 (7.12-12.4)
	FCE 29644A	3.88 (2.90-5.19)
	FCE 26998A*	16.30 (7.18-37.12)
	FCE 29645A	6.58 (4.02-16.97)

(cont..)

Table 1 (cont'd)

Structure	Compound (internal code)	MES-ED ₅₀ (po) mg/Kg
	FCE 28657A*	9.35 (4.34-13.11)
	FCE 29647A	4.34 (3.28-5.75)

where internal code:

- 5 FCE 29088A means 2-(4-(3-fluorobenzyl)oxy)benzylamino)-2-methyl-propanamide, methanesulfonate;
- FCE 26743A* means (S)-2-(4-(3-fluorobenzyl)oxy)benzylamino)propanamide, methanesulfonate;
- FCE 29482A means 2-(4-benzylsulfanyl)benzylamino)-2-methyl-propanamide, methanesulfonate;
- 10 FCE 26727A* means (S)-2-(4-benzylsulfanyl)benzylamino)propanamide, methanesulfonate;
- FCE 29484A means 2-(4-(2-fluorobenzyl)oxy)benzylamino)-2-methyl-propanamide, methanesulfonate;
- FCE 26742A* means (S)-2-(4-(2-fluorobenzyl)oxy)benzylamino)15 propanamide, methanesulfonate;
- FCE 29644A means 2-(4-(3-chlorobenzyl)oxy)benzylamino)-2-methyl-propanamide, methanesulfonate;
- FCE 26193A* means (S)-2-(4-(3-chlorobenzyl)oxy)benzylamino)propanamide, methanesulfonate;
- 20 FCE 29645A means 2-(4-(4-fluorobenzyl)oxy)benzylamino)-2-methyl-propanamide, methanesulfonate;
- FCE 26998A* means (S)-2-(4-(4-fluorobenzyl)oxy)benzylamino)propanamide, methanesulfonate;

-10-

FCF 29647A means 2-(4-(3-fluorobenzyl)oxy)benzylamino)-2-methyl-N-methylpropanamide, methanesulfonate;

FCF 28657A* means (S)-2-(4-(3-fluorobenzyl)oxy)benzylamino)-N-methylpropanamide, methanesulfonate.

5

The comparative activity test data set out in the above table show that the new compounds of the instant invention are from 2 to 4 fold more active than the closely related prior art compounds.

- 10 10 A patient is treated according to the present invention by a method comprising administering to the patient an effective amount of one of the compounds of the invention. In this way the present compounds can be used to treat disorders of the central nervous system, for example epilepsy or Parkinson's
15 disease; or as neuroprotective agents, anti-depressants, hypnotics or anti-spastic agents. The condition of a patient may thus be improved.

20 The compounds of the invention can be administered in a variety of dosage forms, e.g. orally, in the form of tablets, capsules, sugar or film coated tablets, liquid solutions or suspensions; rectally in the form of suppositories; parenterally, e.g. intramuscularly, or by intravenous injection or infusion.

25 The dosage depends on the age, weight, conditions of the patient and on the administration route; for example, the dosage adopted for oral administration to adult humans e.g. for the representative compound of the invention 2-[4-(3-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide may range from about 1 to about 500 mg pro dose, from 1 to 5 times
30 daily.

The invention includes pharmaceutical compositions comprising a compound of the invention, as an active principle, in

association with a pharmaceutically acceptable excipient (which can be a carrier or a diluent).

The pharmaceutical compositions containing the compounds of the invention are usually prepared following conventional 5 methods and are administered in a pharmaceutically suitable form.

For example, the solid oral forms may contain, together with the active compound, diluents, e.g. lactose, destrose, saccharose, cellulose, corn starch or potato starch; 10 lubricants, e.g. silica, talc, stearic acid, magnesium or calcium stearate, and/or polyethylene glycols; binding agents, e.g. starches, arabic gums, gelatin, methylcellulose, carboxymethylcellulose or polyvinyl pyrrolidone; disaggregating agents, e.g. a starch, alginic acid, alginates 15 or sodium starch glycolate; effervescent mixtures; dyestuffs; sweeteners; wetting agents such as lecithin, polysorbates, laurylsulphates; and, in general, non-toxic and pharmacologically inactive substances used in pharmaceutical formulations. Said pharmaceutical preparations may be 20 manufactured in known manner, for example, by means of mixing, granulating, tabletting, sugar-coating, or film-coating processes.

The liquid dispersion for oral administration may be e.g. syrups, emulsions and suspension.

25 The syrups may contain as carrier, for example, saccharose or saccharose with glycerine and/or mannitol and/or sorbitol. The suspension and the emulsion may contain as carrier, for example, a natural gum, agar, sodium alginate, pectin, methylcellulose, carboxymethylcellulose, or polyvinyl alcohol.

30 The suspension or solutions for intramuscular injections may contain, together with the active compound, a

pharmaceutically acceptable carrier, e.g. sterile water, olive oil, ethyl oleate, glycols, e.g. propylene glycol, and, if desired, a suitable amount of lidocaine hydrochloride. The solutions for intravenous injections or infusion may contain 5 as carrier, for example, sterile water or preferably they may be in the form of sterile, aqueous, isotonic saline solutions.

The suppositories may contain together with the active compound a pharmaceutically acceptable carrier, e.g. cocoa 10 butter, polyethylene glycol, a polyoxyethylene sorbitan fatty acid ester surfactant or lecithin.

The following examples illustrate but do not limit the invention.

15 Example 1

2-[4-(3-fluorobenzyl)benzylamino]-2-methyl-propanamide, methanesulfonate (FCE 29088A)

To a solution of 2-amino-2-methylpropanamide, hydrochloride 20 (7.01 g; 0.051 mol) in anhydrous methanol (160 ml), under stirring and nitrogen, 7.0 g of 3Å molecular sieves were added and then, in a single portion, NaBH₃CN (2.31 g; 0.037 mol), after 10 minutes, 10.6 g (0.046 mol) of 4-(3-fluorobenzyl) benzaldehyde were added, in 140 ml of 25 anhydrous methanol. After 24 hours the reaction was completed, the mixture filtered, the solution was evaporated to give a residue which was directly flash-chromatographed on silica gel (eluant: CHCl₃, 98: CH₃OH 2: 30% NH₄OH 0.15) to afford a white solid (6.2 g; 43%). The free base thus 30 obtained was treated with a stoichiometric amount of methanesulfonic acid to yield the title compound (m.p. 209-213°C).

- Analogously, the following products can be obtained, starting from the corresponding aldehyde and the appropriate amide:
- 5 2-[4-(3-chlorobenzylxy)benzylamino]-2-methyl-propanamide,
methanesulfonate, m.p 202-206°C (FCE 29644A);
2-[4-(3-bromobenzylxy)benzylamino]-2-methyl-propanamide,
methanesulfonate, m.p. 197-202°C (FCE 29494A);
2-[4-(4-fluorobenzylxy)benzylamino]-2-methyl-propanamide,
methanesulfonate, m.p. 233°C (FCE 29645A);
10 2-[4-(2-fluorobenzylxy)benzylamino]-2-methyl-propanamide,
methanesulfonate, m.p. 215-220°C (FCE 29484A);
2-[4-(3-fluorobenzylamino)benzylamino]-2-methyl-propanamide,
dihydrochloride, m.p. 165° ca. (dec.) (FCE 29822A);
2-[(4-benzylsulfanyl)benzylamino]-2-methyl-propanamide,
15 methanesulfonate, m.p. 214-215°C (FCE 29482A);
2-[4-(3-fluorobenzylxy)benzylamino]-2-methyl-N-methyl-
propanamide, methanesulfonate, m.p. 213-218°C (FCE 29647A);
and
2- [4-(4-chlorobenzylxy)benzylamino]-2-methyl-propanamide,
20 m.p. 226-227°C (FCE 29485A).

Example 2

2-{[4-(3-fluorobenzylxy)benzyl]methylamino}-2-methyl-
propanamide (FCE 29486)

25 One g. (0.00316 mol) of 2-[(4-(3-fluorobenzylxy)
benzylamino]-2-methyl-propanamide are dissolved in
acetonitrile (50 ml) under a nitrogen stream. To this
mixture, 3.16 ml (0.0389 mol) of 37% formaldehyde and 0.29 g
30 (0.00460 mol) of sodium cyanoborohydride are added at room
temperature. After 20 minutes, glacial acetic acid is dropped

- up to neutrality of the solution. The mixture is stirred for 40 minutes and evaporated to dryness. 40 ml of 2N KOH are added to the residue. After extracting with ethyl acetate, washing with N/2 KOH and then with water and brine, the
5 organic layer is dried on Na₂SO₄, then filtered and evaporated to obtain a residue which is flash-chromatographed on silica gel (eluant: CHCl₃ 200: CH₃OH 3: 30% NH₄OH 0.2) to give 0.75 (72%) of a white solid (m.p. 121-123°C).
- 10 Analogously the following products can be prepared starting from the corresponding secondary amine:
2-{{[4-(3-chlorobenzyl]methoxy}benzyl]methylamino}-2-methylpropanamide; and
2-{{[4-(3-bromobenzyl]methoxy}benzyl]methylamino}-2-methyl-
15 propanamide.

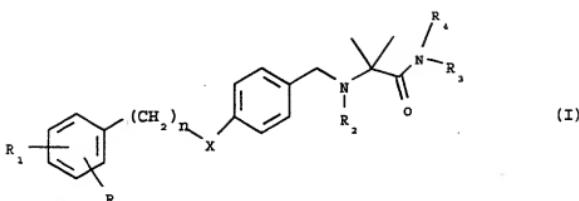
Example 3

With the usual methods of pharmaceutical technique, preparation can be made of capsules having the following
20 composition:

2-[4-(3-fluorobenzyl]methoxy}benzylamino]-2-methyl-	
propanamide, methanesulfonate	50 mg
Talc	2 mg
Corn starch	2 mg
25 Microcristalline cellulose	6 mg
Magnesium stearate	1 mg

CLAIMS

1. A compound of formula (I)



5 wherein:

n is zero, 1, 2 or 3;

X is -O-, -S-, -CH₂- or -NH-;each of R and R₁ independently is hydrogen, C₁-C₆ alkyl, halogen, hydroxy, C₁-C₄ alkoxy or trifluoromethyl;10 each of R₂, R₃ and R₄ independently is hydrogen, C₁-C₆ alkyl or C₃-C₇ cycloalkyl; or a pharmaceutically acceptable salt thereof.

2. A compound of formula (I), according to claim 1,

15 wherein:

n is 1 or 2;

X is -O-, -S- or -NH-;

R is hydrogen;

R₁ is hydrogen or halogen;20 each of R₂, R₃ and R₄ independently is hydrogen or C₁-C₄ alkyl; or a pharmaceutically acceptable salt thereof.

3. A compound of formula (I), according to claim 1,

wherein:

25 n is 1;

X is -O-, -S- or -NH-;

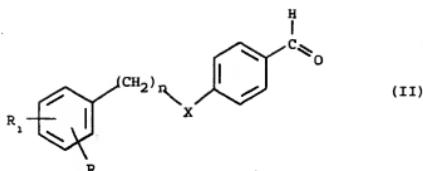
R₁ is hydrogen or halogen;

R, R₂, R₃ and R₄ are hydrogen; or a pharmaceutically acceptable salt thereof.

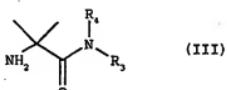
- 5 4. A compound selected from the group consisting of:
2-[4-(3-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;
2-[4-(3-chlorobenzyl)oxy]benzylamino]-2-methyl-propanamide;
2-[4-(4-chlorobenzyl)oxy]benzylamino]-2-methyl-propanamide;
2-[4-(3-bromobenzyl)oxy]benzylamino]-2-methyl-propanamide;
10 2-[4-(4-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;
2-[4-(2-fluorobenzyl)oxy]benzylamino]-2-methyl-propanamide;
2-[4-(3-fluorobenzyl)benzylamino]-2-methyl-propanamide;
2-[4-(benzylsulfanyl)benzylamino]-2-methyl-propanamide;
15 2-[4-(3-fluorobenzyl)benzyl]-methylamino]-2-methyl-
propanamide;
2-[4-(3-fluorobenzyl)benzylamino]-2-methyl-N-methyl-
propanamide;
2-[4-(3-chlorobenzyl)benzyl]methylamino]-2-methyl-
propanamide; and
20 2-[4-(3-bromobenzyl)benzyl]methylamino]-2-methyl-
propanamide, if the case, either as single (S) or (R) isomer
or as a mixture thereof, or a pharmaceutically acceptable
salt thereof.

- 25 5. A process for the preparation of a compound of
formula (I), as defined in claim 1, or a pharmaceutically
acceptable salt thereof, the process comprising:

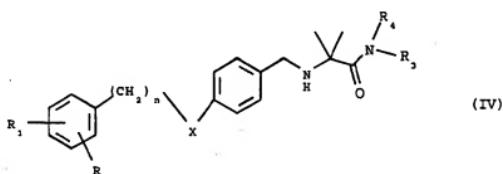
- a) reacting a compound of formula (II)



wherein n, R, R₁ and X are as defined in claim 1, with a compound of formula (III)



- 5 wherein R₃ and R₄ are as defined in claim 1, thus obtaining a compound of formula (I) in which R₂ is hydrogen; or
b) reacting a compound of formula (IV)



- 10 wherein R, R₁, R₃, R₄, n and X are as defined in claim 1, with a compound of formula (V) or (VI)



- wherein W is a halogen atom; R'₂ is a C₁-C₆ alkyl and R''₂ is hydrogen or C₁-C₅ alkyl, thus obtaining a compound of the invention in which R₂ is C₁-C₆ alkyl; and, if desired, 15 converting a compound of the invention into another compound of the invention and/or, if desired, converting a compound of the invention into a pharmaceutically acceptable salt and/or, if desired, converting a salt into a free compound.

- 20 6. A pharmaceutical composition containing a suitable carrier and/or diluent and, as an active principle, a compound of

formula (I), as defined in claim 1, or a pharmaceutically acceptable salt thereof.

7. A compound of formula (I), as defined in claim 1, or
5 a pharmaceutically acceptable salt thereof, for use as an active therapeutic substance.

8. A compound of formula (I), as defined in claim 1, or
10 a pharmaceutically acceptable salt thereof, for use as antiepileptic agent, in the treatment of Parkinson's disease, as neuroprotective agent and in treating and preventing neurodegenerative diseases.

9. A compound of formula (I), as defined in claim 1, or
15 a pharmaceutically acceptable salt thereof, for use in treating ocular damage and retinopathy.

10. A compound of formula (I), as defined in claim 1, or
20 a pharmaceutically acceptable salt thereof, for use as antidepressant, hypnotic antispastic agent.

INTERNATIONAL SEARCH REPORT

National Application No
PCT/EP 96/02961

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C07C237/06 A61K31/16 C07C321/30

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 C07C A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 90 14334 A (ERBA CARLO SPA) 29 November 1990 cited in the application see page 21; claims ---	
A	WO 94 22809 A (ERBA FARMITALIA) 13 October 1994 see claims; examples ---	
A	WO 94 22808 A (ERBA FARMITALIA) 13 October 1994 see claims; examples -----	

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

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- *L" document which may throw doubts on priority claim(s) or which is used to establish the publication date of another citation or other special reason (as specified)
- *O" document referring to an oral disclosure, use, exhibition or other means
- *P" document published prior to the international filing date but later than the priority date claimed

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Date of the actual completion of the international search

22 November 1996

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PCT/EP 96/02961

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